

= s 11 and protecting  
L. 5 FILE USPATFULL  
L3 0 FILE MEDLINE  
L4 2 FILE IFIPAT  
L6 0 FILE USPAT2  
L8 1 FILE CAPLUS  
L7 1 FILE WPIDS  
L8 0 FILE EUROPATFULL  
L9 1 FILE PATOSWO  
L10 1 FILE PCTFULL

TOTAL FOR ALL FILES

L11 11 L1 AND PROTECTING

= dup rem l11

PROCESSING COMPLETED FOR L11

L12 7 DUP REM L11 (4 DUPLICATES REMOVED)

= d l12 1-7 1b1b abs

L12 ANSWER 1 OF 7 USPATFULL

DUPLICATE 1

ACCESSION NUMBER: 2002:126323 USPATFULL

TITLE: **Purification of human troponin I**

INVENTOR(S): Conn, Gregory, Cary, NC, UNITED STATES  
Feardon, Brian, Seattle, WA, UNITED STATES  
Zeng, Xianfang, Northborough, MA, UNITED STATES  
Zhang, Chenming, Blacksburg, VA, UNITED STATES

PATENT ASSIGNEE(S): Biosynth RTP, Inc. (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002064835	A1	20020530
APPLICATION INFO.:	US 2001-903398	A1	20010710 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-217069P	20000710 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	DARBY & DARBY P.C., 805 Third Avenue, New York, NY, 10022	
NUMBER OF CLAIMS:	20	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	11 Drawing Page(s)	
LINE COUNT:	566	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention is directed to methods for purifying **Troponin I**, particularly recombinant Troponin I produced in a bacterial expression system. Recombinant Troponin I can be advantageously purified after reversibly **protecting** the free **sulphydryl** groups, e.g., by forming sulfates. In a specific example, Troponin I reacted with sodium tetrafluoroborate yielded sulfitolyzed Troponin I, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the **sulphydryl** groups yields a highly purified product ready for refolding.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L12 ANSWER 2 OF 7 USPATFULL

DUPLICATE 2

ACCESSION NUMBER: 2002:105940 USPATFULL

TITLE: **Purification of human troponin I**  
 INVENTOR(S): Conn, Gregory, Cary, NC, UNITED STATES  
 Reardon, Brian, Seattle, WA, UNITED STATES  
 Zeng, Xianfang, Northborough, MA, UNITED STATES  
 Zhang, Chenming, Blacksburg, VA, UNITED STATES  
 PATENT ASSIGNEE(S): Diosynth RTP, Inc. (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002055145	A1	20020509
APPLICATION INFO.:	US 2001-998619	A1	20011130 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 2001-903398, filed on 10 Jul 2001, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-217069P	20000710 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	DARBY & DARBY P.C., 805 Third Avenue, New York, NY, 10022	
NUMBER OF CLAIMS:	20	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	11 Drawing Page(s)	
LINE COUNT:	570	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention is directed to methods for purifying **Troponin I**, particularly recombinant Troponin I produced in a bacterial expression system. Recombinant Troponin I can be advantageously purified after reversibly **protecting** the free **sulphydryl** groups, e.g., by forming sulfates. In a specific example, Troponin I reacted with sodium tetrathionate yielded sulfitolyzed Troponin I, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the **sulphydryl** groups yields a highly purified product ready for refolding.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L12 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2002 ACS DUPLICATE 3

ACCESSION NUMBER: 2002:51523 CAPLUS

DOCUMENT NUMBER: 135:101:58

TITLE: Chromatographic **purification** of human **sulphydryl**-protected recombinant **troponin I**

INVENTOR(S): Conn, Gregory; Reardon, Brian; Zeng, Xiangang; Zhang, Chenming

PATENT ASSIGNEE(S): Diosynth RTP, Inc., USA

SOURCE: PCT Int. Appl., 28 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002004512	A2	20020317	WO 2001 US21817	20010710
WO 2002004512	A3	20020516		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CF, CU, CZ, DE, DK, DM, EC, EE, EG, FI, GB, GD, GE, HE, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, ME, NC, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SF, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU,

ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM  
 RW: GH, GM, KE, LS, MW, MG, SD, SL, SS, TZ, UG, ZW, AT, BE, CH, CY,  
 DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,  
 BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

US 2002064835 A1 20020530 US 2001-903398 20010710  
 US 2002055145 A1 20020509 US 2001-998619 20011130

PRIORITY APPLN. INFO.: US 2000-217069P P 20000710  
 US 2001-903398 A1 20010710

AB The invention is directed to methods for purifying **troponin I**, particularly recombinant **troponin I** produced in a bacterial expression system. Recombinant **troponin I** can be advantageously purified after reversibly **protecting** the free **sulphydryl** groups, e.g. by forming sulfates. In a specific example, **troponin I** reacted with sodium tetrathionate yielded sulfitolysed **troponin I**, which was purified by chromatog. on an anion exchanger, followed by hydrophobic interaction chromatog. Facile deprotection of the **sulphydryl** groups yields a highly purified product ready for refolding.

L12 ANSWER 4 OF 7 USPATFULL

ACCESSION NUMBER: 2002:85170 USPATFULL  
 TITLE: Neuropeptide-like polypeptide zpep17  
 INVENTOR(S): Sheppard, Paul D., Granite Falls, WA, UNITED STATES  
 Bishop, Paul D., Fall City, WA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002045210	A1	20020418
APPLICATION INFO.:	US 2001-776795	A1	20010305 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-180314P	20000304 (60)
	US 2000-180896P	20000307 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Jennifer K. Johnson, ZymoGenetics, Inc, 1201 Eastlake Avenue East, Seattle, WA, 98102	
NUMBER OF CLAIMS:	26	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	12 Drawing Page(s)	
LINE COUNT:	4459	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to polynucleotide and polypeptide molecules for zpep17, a novel secreted protein. The polynucleotides encoding zpep17, may, for example, be used to identify a region of the genome associated with human disease states. The present invention also includes methods for producing the protein, uses therefor and antibodies thereto.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L12 ANSWER 5 OF 7 USPATFULL

ACCESSION NUMBER: 2002:43270 USPATFULL  
 TITLE: Methods for analyzing protein binding events  
 INVENTOR(S): Hefti, John J., San Francisco, CA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002028461	A1	20020307
APPLICATION INFO.:	US 2001-913474	A1	20010806 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1999-365 80, filed on 2 Aug 1999, GRANTED, Pat. No. US 6287374 Continuation-in-part of Ser. No. US 1999-243124, filed on 1 Feb 1999,		

PENDING

	NUMBER	DATE
PRIORITY INFORMATION:	US 1993-73445P	19980202 (60)
	US 1993-134740P	19990518 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Richard L. Neeley, Clifford B. Perry, Signature	
	BioScience, Inc., 21124 Cabot Boulevard, Hayward, CA,	
	94545-1130	
NUMBER OF CLAIMS:	45	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	37 Drawing Page(s)	
LINE COUNT:	4041	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a variety of methods of analyzing protein binding events using a system capable of directly detecting protein/ligand complexes based upon the dielectric properties of the complex. The system can be used in a variety of analyses involving protein binding events, such as screening ligand libraries, characterizing protein binding interactions, and identifying ligands. The system can also be utilized in diverse analytical and diagnostic applications.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L12 ANSWER 6 OF 7 PCTFULL COPYRIGHT 2002 Univentio  
ACCESSION NUMBER: 2002036624 PCTFULL ED 20020523 EW 200219  
TITLE (ENGLISH): METHODS AND COMPOSITIONS RELATING TO FORTILIN, AN  
ANTI-APOPTOTIC MOLECULE, AND MODULATORS OF FORTILIN  
TITLE (FRENCH): PROCEDES ET COMPOSITIONS ASSOCIES A LA FORTILINE, UNE  
MOLECULE ANTI-APOPTOTIQUE, ET MODULATEURS DE FORTILINE  
INVENTOR(S): FUJISE, Kenichi; YEH, Edward  
PATENT ASSIGNEE(S): BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM, for  
all designates States except US; FUJISE, Kenichi, for  
US only; YEH, Edward, for US only  
AGENT: SHISHIMA, Gina, N.  
LANGUAGE OF PUBL.: English  
LANGUAGE OF FILING: English  
DOCUMENT TYPE: Patent  
PATENT INFORMATION:

	NUMBER	KIND	DATE
DESIGNATED STATES:	WO 2002036624	A2	20020510
	AE AG AL AM AT AU AZ BA BB BG BR BY BE CA CH CN CO CR		
	CU CZ DE DK DM DZ EC EE ES FI GR GD GE GH GM HR HU ID		
	IL IN IS JP KE KG KP KR KZ LC LK LF LS LT LU LV MA MD		
	ME MK MN MW MX MZ NO NZ PH PL PT RO RU SD SE SG SI SK		
	SL TM TN TR TT TZ UA UG US UZ VN YU ZA ZW ZY GM KE LS		
	MW MZ SD SL SZ TZ UG ZW AM AZ BY EG KE MD EU TJ TM AT		
	BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR		
	BF BI CF CG CI CM GA GN GQ GW ML MF NE SN TD TG		

APPLICATION INFO.: WO 2001-US42985 A 20011030  
PRIORITY INFO.: US 2000-60/244,416 20001030

ABEN The polypeptide Fortilin (also known as Translationally Controlled Tumour Protein, TTP) specifically interacts with p53, a tumor suppressor involved in the induction of apoptosis and the normal growth regulation of a cell. Fortilin also specifically binds MLL1 (Myeloid Cell Leukemia 1). Fortilin has the ability to prevent apoptosis, which may be unregulated in hyperproliferative cells. The present invention is directed at compositions and methods involving a Fortilin modulator, which can induce apoptosis, for the prevention, treatment, or diagnosis of hyperproliferative diseases and conditions, including cancer and atherosclerosis. It is directed also at compositions and methods

involving Fortilin, which can inhibit apoptosis, for the treatment of diseases and condition characterized by apoptosis, including certain vascular conditions.

ABFR Le polypeptide fortiline (egalement appele proteine tumorale de regulation de traduction, TCTP) interagit specifiquement avec p53, un supprimeur de tumeur intervenant dans l'induction de l'apoptose et la regulation de la croissance normale d'une cellule. La fortiline se lie aussi specifiquement a MCL1 (leucemie myeloide 1). La fortiline est capable de prevenir l'apoptose, qui peut etre dereglee dans des cellules hyperproliferatives. L'invention concerne des compositions et des procedes comprenant un modulateur de fortiline, capable d'induire l'apoptose, pour prevenir, traiter ou diagnostiquer des maladies ou des affections hyperproliferatives, y compris le cancer et l'atherosclerose ; ainsi que des compositions et des procedes comprenant la fortiline, capable d'inhiber l'apoptose, pour traiter des maladies et affections caracterisees par l'apoptose, y compris certaines affections vasculaires.

L12 ANSWER 7 OF 7 USPATFULL

ACCESSION NUMBER: 2001:152781 USPATFULL  
TITLE: Methods for analyzing protein binding events  
INVENTOR(S): Hefti, John, San Francisco, CA, United States  
PATENT ASSIGNEE(S): Signature BioScience, Inc., Hayward, CA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6287874	B1	20010911
APPLICATION INFO.:	US 1999-365580		19990802 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1999-243194, filed on 1 Feb 1999		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1998-73445P	19980202 (60)
	US 1999-134740P	19990518 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Horlick, Kenneth R.	
ASSISTANT EXAMINER:	Strzelecka, Teresa	
LEGAL REPRESENTATIVE:	Ausenhuss, Scott L., Perry, Clifford B., Neeley, Richard L.	
NUMBER OF CLAIMS:	45	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	44 Drawing Figure(s); 33 Drawing Page(s)	
LINE COUNT:	4099	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a variety of methods of analyzing protein binding events using a system capable of directly detecting protein/ligand complexes based upon the dielectric properties of the complex. The system can be used in a variety of analyses involving protein binding events, such as screening ligand libraries, characterizing protein binding interactions, and identifying ligands. The system can also be utilized in diverse analytical and diagnostic applications.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

90 s troponin and purification

L13 270 FILE USPATFULL  
L14 479 FILE MEDLINE  
L15 5 FILE IFIPAT  
L16 5 FILE USPAT2  
L17 204 FILE CAPLUS

L18 11 FILE WPIDS  
 L19 45 FILE EUROPATFULL  
 L20 4 FILE PATOSWO  
 L21 19 FILE PCTFULL

TOTAL FOR ALL FILES

L32 1042 TROPONIN AND PURIFICATION

=> s l32 and (sulfhydryl (w) group)

L33 35 FILE USPATFULL  
 L34 3 FILE MEDLINE  
 L35 2 FILE IFIPAT  
 L36 2 FILE USPATE  
 L37 1 FILE CAPLUS  
 L38 1 FILE WPIDS  
 L39 1 FILE EUROPATFULL  
 L30 1 FILE PATOSWO  
 L31 1 FILE PCTFULL

TOTAL FOR ALL FILES

L33 47 L32 AND (SULFHYDRYL (W) GROUP)

=> dup rem l33

PROCESSING COMPLETED FOR L32

L33 41 DUP REM L32 (6 DUPLICATES REMOVED)

=> d l33 1-41 ibib abs

L33 ANSWER 1 OF 41 USPATFULL DUPLICATE 1  
 ACCESSION NUMBER: 2002:126323 USPATFULL  
 TITLE: **Purification of human troponin I**  
 INVENTOR(S): Conn, Gregory, Cary, NC, UNITED STATES  
 Reardon, Brian, Seattle, WA, UNITED STATES  
 Zeng, Xianfang, Northborough, MA, UNITED STATES  
 Zhang, Chenming, Blacksburg, VA, UNITED STATES  
 PATENT ASSIGNEE(S): Diosynth ETP, Inc. (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002064835	A1	20020530
APPLICATION INFO.:	US 2001-963338	A1	20010710 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-217069P	20000710 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	DARBY & DARBY P.C., 805 Third Avenue, New York, NY, 10022	
NUMBER OF CLAIMS:	20	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	11 Drawing Page(s)	
LINE COUNT:	566	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention is directed to methods for purifying **Troponin I**, particularly recombinant Troponin I produced in a bacterial expression system. Recombinant Troponin I can be advantageously purified after reversibly protecting the free **sulfhydryl groups**, e.g., by forming sulfates. In a specific example, Troponin I reacted with sodium tetrathionate yielded Sulfatolyzed Troponin I, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the **sulfhydryl groups** yields a highly purified product ready for refolding.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 2 OF 41 USPATFULL

DUPLICATE 2

ACCESSION NUMBER: 2002:105940 USPATFULL

TITLE: **Purification of human troponin I**

INVENTOR(S): Conn, Gregory, Cary, NC, UNITED STATES  
Reardon, Brian, Seattle, WA, UNITED STATES  
Zeng, Xianfang, Northborough, MA, UNITED STATES  
Zhang, Chenming, Blacksburg, VA, UNITED STATES

PATENT ASSIGNEE(S): Biosynth RTP, Inc. (U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 2002:1055145 A1 20020509  
APPLICATION INFO.: US 2001-998619 A1 20011130 (9)  
RELATED APPLN. INFO.: Continuation of Ser. No. US 2001-903398, filed on 10  
Jul 2001, PENDING

NUMBER DATE

PRIORITY INFORMATION: US 2000-217069P 20000710 (69)  
DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: LARBY & LARBY P.C., 805 Third Avenue, New York, NY,  
10022  
NUMBER OF CLAIMS: 20  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 11 Drawing Page(s)  
LINE COUNT: 570

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention is directed to methods for purifying **Troponin I**, particularly recombinant Troponin I produced in a bacterial expression system. Recombinant Troponin I can be advantageously purified after reversibly protecting the free **sulfhydryl groups**, e.g., by forming sulfates. In a specific example, Troponin I reacted with sodium tetrathionate yielded sulfitylized Troponin I, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the **sulfhydryl groups** yields a highly purified product ready for refolding.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 3 OF 41 USPATFULL

DUPLICATE 3

ACCESSION NUMBER: 2002:16843 USPATFULL

TITLE: **RESONANT BIO-ASSAY DEVICE AND TEST SYSTEM FOR DETECTING MOLECULAR-BINDING EVENTS**

INVENTOR(S): HEFTI, JOHN, SAN FRANCISCO, CA, UNITED STATES

NUMBER KIND DATE

PATENT INFORMATION: US 2002009723 A1 20020124  
US 6376258 B2 20020423  
APPLICATION INFO.: US 2000-489846 A1 20000110 (9)  
RELATED APPLN. INFO.: Continuation of Ser. No. US 1999-365578, filed on 2 Aug  
1999, PENDING Continuation-in-part of Ser. No. US  
1999-243196, filed on 1 Feb 1999, PENDING  
Continuation-in-part of Ser. No. US 1999-243194, filed  
on 1 Feb 1999, PENDING

NUMBER DATE

PRIORITY INFORMATION: US 1998-72445P 19980202 (60)  
DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: SIGNATURE BIOSCIENCE, INC., 21124 CABOT BLVD., HAYWARD,  
CA, 94545-1130  
NUMBER OF CLAIMS: 64  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 38 Drawing Page(s)  
LINE COUNT: 3548  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Systems and methods are presented for detecting molecular binding events and other environmental effects using the unique dielectric properties of the bound molecular structure or structures. A molecular binding region is coupled along the surface of a signal path. A test signal is propagated along the signal path, whereby the test signal couples to the molecular binding region, and in response, exhibits a signal response.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 4 OF 41 USPATFULL DUPLICATE 4  
ACCESSION NUMBER: 2002:122429 USPATFULL  
TITLE: Computer program and database structure for detecting molecular binding events  
INVENTOR(S): Hefti, John, San Francisco, CA, United States  
PATENT ASSIGNEE(S): Signature BioScience, Inc., Hayward, CA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6395480	B1	20020528
	US 2002072857	A1	20020613
APPLICATION INFO.:	US 1999-243196		19990201 (9)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	GRANTED		
PRIMARY EXAMINER:	Brusca, John S.		
ASSISTANT EXAMINER:	Kim, Young		
LEGAL REPRESENTATIVE:	Perry, Clifford B.		
NUMBER OF CLAIMS:	21		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	51 Drawing Figure(s); 28 Drawing Page(s)		
LINE COUNT:	3363		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Systems and methods for detecting molecular binding events and other environmental effects using the unique dielectric properties of the bound molecular structure or structures are presented. A molecular binding layer is coupled along the surface of a signal path. A test signal is propagated along the signal path, whereby the test signal couples to the molecular binding layer, and in response, exhibits a signal response.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 5 OF 41 CAPLUS COPYRIGHT 2002 ACS DUPLICATE 5  
ACCESSION NUMBER: 2002:51573 CAPLUS  
DOCUMENT NUMBER: 136:101258  
TITLE: Chromatographic **purification** of human  
sulfhydryl-protected recombinant **troponin I**  
INVENTOR(S): Conn, Gregory; Reardon, Brian; Zeng, Xiangang; Zhang, Chenming  
PATENT ASSIGNEE(S): Diosynth FTF, Inc., USA  
SOURCE: PCT Int. Appl., 28 pp.  
COMMEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 2002004512 A2 20020117 WO 2001-US21817 20010710  
WO 2002004512 A1 20020516

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,  
CO, CR, CU, CZ, DE, DK, DM, EC, EE, ES, FI, GB, GD, GE, HR, HU,  
ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,  
LV, MA, MD, MG, MK, MN, MW, MX, ME, NO, NZ, PL, PT, RO, RU, SD,  
SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU,  
ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,  
DE, DK, ES, FI, FR, GB, GR, IE, IT, LJ, MC, NL, PT, SE, TR, BF,  
BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

US 2002064835 A1 20020530 US 2001-903398 20010710  
US 2002055145 A1 20020509 US 2001-998619 20011130

PRIORITY APPLN. INFO.: US 2000-217069P P 20000710  
US 2001-903398 A1 20010710

AB The invention is directed to methods for purifying **troponin I**, particularly recombinant **troponin I** produced in a bacterial expression system. Recombinant **troponin I** can be advantageously purified after reversibly protecting the free **sulfhydryl groups**, e.g. by forming sulfates. In a specific example, **troponin I** reacted with sodium tetrathionate yielded sulfitolyzed **troponin I**, which was purified by chromatog. on an anion exchanger, followed by hydrophobic interaction chromatog. Facile deprotection of the **sulfhydryl groups** yields a highly purified product ready for refolding.

L33 ANSWER 6 OF 41 USPTAFULL

ACCESSION NUMBER: 2002:149299 USPTAFULL  
TITLE: Death domain-containing receptor polynucleotides, polypeptides, and antibodies  
INVENTOR(S): Ni, Jian, Germantown, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002077458	A1	20020620
APPLICATION INFO.:	US 2001-835788	A1	20010417 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US8666, filed on 17 Oct 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-159585P	19991018 (60)
	US 1999-167246P	19991124 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	12	
EXEMPLARY CLAIM:	1	
LINE COUNT:	14143	

AB The present invention relates to novel human BDCP polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human BDCP polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human BDCP polypeptides.

L33 ANSWER 7 OF 41 USPTAFULL

ACCESSION NUMBER: 2002:149131 USPTAFULL  
TITLE: 18 human secreted proteins  
INVENTOR(S): Ruben, Steven M., Olney, MD, UNITED STATES

Rosen, Craig A., Laytonsville, MD, UNITED STATES  
 Li, Yi, Sunnyvale, CA, UNITED STATES  
 Zeng, Zhizhen, Lansdale, PA, UNITED STATES  
 Kyaw, Hla, Frederick, MD, UNITED STATES  
 Fischer, Carrie L., Burke, VA, UNITED STATES  
 Li, Haodong, Gaithersburg, MD, UNITED STATES  
 Seppet, Daniel R., Centreville, VA, UNITED STATES  
 Gentz, Reiner L., Rockville, MD, UNITED STATES  
 Wei, Ying-Fei, Berkeley, CA, UNITED STATES  
 Moore, Paul A., Germantown, MD, UNITED STATES  
 Young, Paul E., Gaithersburg, MD, UNITED STATES  
 Greene, John M., Gaithersburg, MD, UNITED STATES  
 Ferrie, Ann M., Tewksbury, MA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002077287	A1	20020620
APPLICATION INFO.:	US 2001-853659	A1	20010511 (9)
RELATED APPLN. INFO.:	Continuation in-part of Ser. No. US 1998-152060, filed on 11 Sep 1998, UNKNOWN		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	APPLICATION		
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850		
NUMBER OF CLAIMS:	23		
EXEMPLARY CLAIM:	1		
LINE COUNT:	17779		

AB The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

L33 ANSWER 8 OF 41 USPATFULL

ACCESSION NUMBER: 2002:149114 USPATFULL  
 TITLE: Nucleic acids, proteins, and antibodies  
 INVENTOR(S): Fosen, Craig A., Laytonsville, MD, UNITED STATES  
 Fuben, Steven M., Olney, MD, UNITED STATES  
 Barash, Steven C., Rockville, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002077270	A1	20020620
APPLICATION INFO.:	US 2001-764848	A1	20010117 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-172065P	20000131 (60)
	US 2000-180638P	20000204 (60)
	US 2000-214886P	20000628 (60)
	US 2000-217487P	20000711 (60)
	US 2000-225758P	20000814 (60)
	US 2000-220963P	20000726 (60)
	US 2000-217496P	20000711 (60)
	US 2000-216447P	20000814 (60)
	US 2000-218200P	20000714 (60)
	US 2000-225757P	20000814 (60)
	US 2000-216808P	20000822 (60)
	US 2000-216647P	20000707 (60)
	US 2000-225260P	20000814 (60)
	US 2000-216880P	20000707 (60)
	US 2000-225270P	20000814 (60)

US 2000-251869P	20001208 (60)
US 2000-235834P	20000927 (60)
US 2000-234274P	20000921 (60)
US 2000-234233P	20000921 (60)
US 2000-238924P	20000830 (60)
US 2000-224518P	20000814 (60)
US 2000-236369P	20000929 (60)
US 2000-224519P	20000814 (60)
US 2000-220964P	20000726 (60)
US 2000-241809P	20001020 (60)
US 2000-249299P	20001117 (60)
US 2000-236327P	20000929 (60)
US 2000-241785P	20001020 (60)
US 2000-244617P	20001101 (60)
US 2000-225268P	20000814 (60)
US 2000-236368P	20000929 (60)
US 2000-251856P	20001208 (60)
US 2000-251868P	20001208 (60)
US 2000-229344P	20000901 (60)
US 2000-234997P	20000925 (60)
US 2000-229343P	20000901 (60)
US 2000-229345P	20000901 (60)
US 2000-229287P	20000901 (60)
US 2000-229513P	20000905 (60)
US 2000-231413P	20000908 (60)
US 2000-229509P	20000905 (60)
US 2000-236367P	20000929 (60)
US 2000-237039P	20001002 (60)
US 2000-237038P	20001002 (60)
US 2000-236370P	20000929 (60)
US 2000-236802P	20001002 (60)
US 2000-237037P	20001002 (60)
US 2000-237040P	20001002 (60)
US 2000-240960P	20001020 (60)
US 2000-239935P	20001013 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
ROCKVILLE, MD, 20850  
NUMBER OF CLAIMS: 24  
EXEMPLARY CLAIM: 1  
LINE COUNT: 20057

AB The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

L33 ANSWER 9 OF 41 USPATEL

ACCESSION NUMBER: 2002:143614 USPATEL  
TITLE: 28 Human secreted proteins  
INVENTOR(S): Ruben, Steven M., Olney, MD, UNITED STATES  
Rosen, Craig A., Laytonville, MD, UNITED STATES  
Li, Yi, Sunnyvale, CA, UNITED STATES  
Geng, ShiZhen, Lansdale, PA, UNITED STATES  
Kyaw, Hla, Frederick, MD, UNITED STATES

Fischer, Carrie L., Burke, VA, UNITED STATES  
 Li, Haodong, Gaithersburg, MD, UNITED STATES  
 Soppet, Daniel R., Centreville, VA, UNITED STATES  
 Gentz, Reiner L., Rockville, MD, UNITED STATES  
 Wei, Ying-Fei, Berkeley, CA, UNITED STATES  
 Moore, Paul A., Germantown, MD, UNITED STATES  
 Young, Paul E., Gaithersburg, MD, UNITED STATES  
 Greene, John M., Gaithersburg, MD, UNITED STATES  
 Ferrie, Ann M., Painted Post, NY, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002076756	A1	20020620
APPLICATION INFO.:	US 2001-853161	A1	20010511 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2001-265583P	20010202 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	23	
EXEMPLARY CLAIM:	1	
LINE COUNT:	17788	

AB The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

L33 ANSWER 10 OF 41 USPTAFULL

ACCESSION NUMBER: 2001:141609 USPTAFULL  
 TITLE: Transferrin polynucleotides, polypeptides, and antibodies  
 INVENTOR(S): Fuben, Steven M., Olney, MD, UNITED STATES  
 Shi, Yangu, Gaithersburg, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002076756	A1	20020613
APPLICATION INFO.:	US 2001-891126	A1	20010626 (9)
RELATED APPLN. INFO.:	Continuation-in part of Ser. No. WD 2000-US34769, filed on 21 Dec 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-171595P	19991213 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	22	
EXEMPLARY CLAIM:	1	
LINE COUNT:	12048	

AB The present invention relates to novel human transferrin polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human transferrin polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human transferrin polypeptides.

L33 ANSWER 11 OF 41 USPATFULL

ACCESSION NUMBER: 2002:133469 USPATFULL

TITLE: Serine protease polynucleotides, polypeptides, and antibodies

INVENTOR(S): Shi, Yanqun, Gaithersburg, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES  
Ni, Jian, Germantown, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002068420	A1	20020506
APPLICATION INFO.:	US 2001-804156	A1	20010313 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-189025P	20000314 (50)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	22	
EXEMPLARY CLAIM:	1	
LINE COUNT:	13119	

AB The present invention relates to novel human serine protease polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human serine protease polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human serine protease polypeptides.

L33 ANSWER 12 OF 41 USPATFULL

ACCESSION NUMBER: 2002:136703 USPATFULL

TITLE: Immunoglobulin superfamily polynucleotides, polypeptides, and antibodies

INVENTOR(S): Young, Paul E., Gaithersburg, MD, UNITED STATES  
Ni, Jian, Rockville, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES  
Shi, Yanqun, Gaithersburg, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002068420	A1	20020530
APPLICATION INFO.:	US 2001-799514	A1	20010307 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US23662, filed on 29 Aug 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-152248P	19990903 (50)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	22	
EXEMPLARY CLAIM:	1	
LINE COUNT:	12447	

AB The present invention relates to novel human Ig like polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human Ig-like polypeptides. The invention further relates to diagnostic and

therapeutic methods useful for diagnosing and treating disorders related to these novel human Ig-like polypeptides.

L33 ANSWER 13 OF 41 USPTFULL

ACCESSION NUMBER: 2002:126332 USPTFULL  
TITLE: Human protein tyrosine phosphatase polynucleotides, polypeptides, and antibodies  
INVENTOR(S): Shi, Yangu, Gaithersburg, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002064844	A1	20020530
APPLICATION INFO.:	US 2001-906779	A1	20010718 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2001-US1563, filed on 17 Jan 2001, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-176306P	20000118 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	22	
EXEMPLARY CLAIM:	1	
LINE COUNT:	12129	

AB The present invention relates to novel human PTPase polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human PTPase polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human PTPase polypeptides.

L33 ANSWER 14 OF 41 USPTFULL

ACCESSION NUMBER: 2002:126317 USPTFULL  
TITLE: Human tumor necrosis factor delta and epsilon  
INVENTOR(S): Yu, Guo-Liang, Berkeley, CA, UNITED STATES  
Ni, Jian, Germantown, MD, UNITED STATES  
Gentz, Eriener L., Rockville, MD, UNITED STATES  
Billon, Patrick J., Carlsbad, CA, UNITED STATES  
PATENT ASSIGNEE(S): Human Genome Sciences, Inc., Rockville, MD, UNITED STATES, 20850 (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002064829	A1	20020530
APPLICATION INFO.:	US 2001-879919	A1	20010614 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1997-815783, filed on 12 Mar 1997, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1996-16812P	19960314 (60)
	US 2001-123499P	20010625 (60)
	US 2001-177478P	20010423 (60)
	US 2001-176143P	20010316 (60)
	US 2001-154875P	20010213 (60)
	US 2001-141952P	20010223 (60)
	US 2000-211537P	20000615 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: 63

EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 11 Drawing Page(s)

LINE COUNT: 13531

AB The invention relates to human TNF delta and TNF epsilon polypeptides, polynucleotides encoding the polypeptides, methods for producing the polypeptides, in particular by expressing the polynucleotides, and agonists and antagonists of the polypeptides. The invention further relates to methods for utilizing such polynucleotides, polypeptides, agonists and antagonists for applications, which relate, in part, to research, diagnostic and clinical arts.

L33 ANSWER 15 OF 41 USPTAFULL

ACCESSION NUMBER: 2002:126314 USPTAFULL

TITLE: Cytokine receptor-like polynucleotides, polypeptides, and antibodies

INVENTOR(S): Fuben, Steven M., Olney, MD, UNITED STATES

Ni, Jian, Germantown, MD, UNITED STATES

Young, Paul E., Gaithersburg, MD, UNITED STATES

Shi, Yanggu, Gaithersburg, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002064826	A1	20020530
APPLICATION INFO.:	US 2001-874069	A1	20010606 (9)
RELATED APPLN. INFO.:	Continuation in-part of Ser. No. WO 2000-US32525, filed on 30 Nov 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999 168621P	19991203 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: 22

EXEMPLARY CLAIM: 1

LINE COUNT: 12089

AB The present invention relates to novel human cytokine receptor-like polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human cytokine receptor-like polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human cytokine receptor-like polypeptides.

L33 ANSWER 16 OF 41 USPTAFULL

ACCESSION NUMBER: 2002:126306 USPTAFULL

TITLE: 30 human secreted proteins

INVENTOR(S): Ni, Jian, Germantown, MD, UNITED STATES

Baker, Kevin P., Darnestown, MD, UNITED STATES

Birso, Charles E., North Potomac, MD, UNITED STATES

Esposito, Michele, Bethesda, MD, UNITED STATES

Ematsoulis, George A., Silver Spring, MD, UNITED STATES

Fisen, Craig A., Laytonsville, MD, UNITED STATES

Jeppet, Daniel E., Centreville, VA, UNITED STATES

Young, Paul E., Gaithersburg, MD, UNITED STATES

Biner, Bernhard, Gaithersburg, MD, UNITED STATES

Juan, D. Roxanne, Bethesda, MD, UNITED STATES

Olsen, Henrik S., Gaithersburg, MD, UNITED STATES

LaFleur, David W., Washington, DC, UNITED STATES  
 Moore, Paul A., Germantown, MD, UNITED STATES  
 Shi, Yanggu, Gaithersburg, MD, UNITED STATES  
 Wei, Ping, Brookeville, MD, UNITED STATES  
 Florence, Kimberly A., Rockville, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002064818	A1	20020530
APPLICATION INFO.:	US 2001-789561	A1	20010222 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US24008, filed on 31 Aug 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-152317P	19990903 (60)
	US 1999-152315P	19990903 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	23	
EXEMPLARY CLAIM:	1	
LINE COUNT:	24623	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human secreted proteins and isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 17 OF 41 USPATFULL

ACCESSION NUMBER: 2002:119538 USPATFULL  
 TITLE: Nucleic acids, proteins, and antibodies  
 INVENTOR(S): Eosen, Craig A., Laytonsville, MD, UNITED STATES  
 Fuher, Steven M., Olney, MD, UNITED STATES  
 Barash, Steven C., Rockville, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002061521	A1	20020523
APPLICATION INFO.:	US 2001-764869	A1	20010117 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-179065P	20000131 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	14	
EXEMPLARY CLAIM:	1	
LINE COUNT:	7727	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel cardiovascular system related polynucleotides and the polypeptides encoded by these polynucleotides herein collectively known as "cardiovascular system antigens," and the use of such cardiovascular system antigens for detecting disorders of the cardiovascular system, particularly the presence of cancer of cardiovascular system tissues and cancer metastases. More specifically, isolated cardiovascular system associated nucleic acid molecules are



provided encoding novel cardiovascular system associated polypeptides. Novel cardiovascular system polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human cardiovascular system associated polynucleotides and/or polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to the cardiovascular system, including cancer of cardiovascular system tissues, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting the production and function of the polypeptides of the present invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 18 OF 41 USPATFULL

ACCESSION NUMBER: 2002:105937 USPATFULL

TITLE: Major intrinsic protein (MIP) like polynucleotides, polypeptides, and antibodies

INVENTOR(S): Ruben, Steven A., Olney, MD, UNITED STATES  
Ni, Jian, Germantown, MD, UNITED STATES

PATENT ASSIGNEE(S): Human Genome Sciences, Inc., Rockville, MD (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002055142	A1	20020509
APPLICATION INFO.:	US 2001-862419	A1	20010523 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US31919, filed on 21 Nov 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-167247F	19991124 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	22	
EXEMPLARY CLAIM:	1	
LINE COUNT:	11747	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human MIP-like polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human MIP-like polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human MIP-like polypeptides.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 19 OF 41 USPATFULL

ACCESSION NUMBER: 2002:92088 USPATFULL

TITLE: Eriptide domain-containing polynucleotides, polypeptides, and antibodies

INVENTOR(S): Ni, Jian, Germantown, MD, UNITED STATES  
Moore, Paul A., Germantown, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002051484	A1	20020502

APPLICATION INFO.: US 2001-848288 A1 20010504 (9)  
 RELATED APPLN. INFO.: Continuation-in-part of Ser. No. WO 2000-US30664, filed  
 on 8 Nov 2000, UNKNOWN

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-164853P	19991112 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	33	
EXEMPLARY CLAIM:	1	
LINE COUNT:	12041	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human KDC polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human KDC polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human KDC polypeptides.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 20 OF 41 USPTFULL

ACCESSION NUMBER: 2002:85190 USPTFULL  
 TITLE: Nucleic acids, proteins, and antibodies  
 INVENTOR(S): Rosen, Craig A., Laytonsville, MD, UNITED STATES  
 Rubin, Steven M., Olney, MD, UNITED STATES  
 Barash, Steven C., Rockville, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002045230	A1	20020418
APPLICATION INFO.:	US 2001-908711	A1	20010720 (9)
RELATED APPLN. INFO.:	Continuation-in part of Ser. No. WO 2001-US1360, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764867, filed on 17 Jan 2001, UNKNOWN Continuation-in part of Ser. No. WO 2001-US1344, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764892, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. WO 2001-US1345, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764888, filed on 17 Jan 2001, UNKNOWN Continuation-in part of Ser. No. WO 2001-US1329, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764905, filed on 17 Jan 2001, UNKNOWN Continuation-in part of Ser. No. US 2001-764891, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. WO 2001-US1339, filed on 17 Jan 2001, UNKNOWN Continuation-in part of Ser. No. US 2001-764899, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. WO 2001-US1340, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764874, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. WO 2001-US1334, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764898, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. WO 2001-US1320, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764893, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. US 2001-764902, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser. No. WO 2001-US1289, filed on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser.		

No. US 2001-764870, filed on 17 Jan 2001, UNKNOWN  
Continuation-in-part of Ser. No. WO 2001-US1348, filed  
on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser.  
No. US 2001-764882, filed on 17 Jan 2001, UNKNOWN  
Continuation-in-part of Ser. No. WO 2001-US1347, filed  
on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser.  
No. US 2001-764896, filed on 17 Jan 2001, UNKNOWN  
Continuation-in-part of Ser. No. WO 2001-US1307, filed  
on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser.  
No. US 2001-764864, filed on 17 Jan 2001, UNKNOWN  
Continuation-in-part of Ser. No. WO 2001-US1341, filed  
on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser.  
No. US 2001-764856, filed on 17 Jan 2001, UNKNOWN  
Continuation-in-part of Ser. No. WO 2001-US1336, filed  
on 17 Jan 2001, UNKNOWN Continuation-in-part of Ser.  
No. US 2001-764868, filed on 17 Jan 2001, UNKNOWN  
Continuation-in-part of Ser. No. WO 2001-US1312, filed  
on 17 Jan 2001, UNKNOWN

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-179066P	20000131 (60)
	US 2000-180628P	20000204 (60)
	US 2000-251868P	20001209 (60)
	US 2000-232398P	20000914 (60)
	US 2000-249300P	20001117 (60)
	US 2000-251990P	20001208 (60)
	US 2000-250160P	20001201 (60)
	US 2000-209467P	20000607 (60)
	US 2000-179065P	20000131 (60)
	US 2000-180628P	20000204 (60)
	US 2000-214886P	20000628 (60)
	US 2000-217487P	20000711 (60)
	US 2000-225758P	20000814 (60)
	US 2000-220963P	20000726 (60)
	US 2000-217496P	20000711 (60)
	US 2000-225447P	20000814 (60)
	US 2000-218290P	20000714 (60)
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	US 2000-226868P	20000822 (60)
	US 2000-216647P	20000707 (60)
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	US 2000-216880P	20000707 (60)
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	US 2000-239834P	20000927 (60)
	US 2000-234274P	20000921 (60)
	US 2000-234273P	20000921 (60)
	US 2000-238224P	20000830 (60)
	US 2000-224518P	20000814 (60)
	US 2000-236269P	20000929 (60)
	US 2000-234519P	20000814 (60)
	US 2000-239964P	20000726 (60)
	US 2000-241309P	20001020 (60)
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	US 2000-244627P	20001101 (60)
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	US 2000-231856P	20001108 (60)
	US 2000-231858P	20001208 (60)
	US 2000-239344P	20000901 (60)
	US 2000-234347P	20000925 (60)
	US 2000-229343P	20000901 (60)

US 2000-249300P	20001117 (60)
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US 2000-246610P	20001108 (60)
US 2000-246611P	20001108 (60)
US 2000-230437P	20000906 (60)
US 2000-251990P	20001203 (60)
US 2000-251988P	20001205 (60)
US 2000-251930P	20001205 (60)
US 2000-251479P	20001206 (60)
US 2000-256719P	20001206 (60)
US 2000-250160P	20001201 (60)
US 2000-251989P	20001208 (60)
US 2000-250391P	20001201 (60)
US 2000-254097P	20001211 (60)
US 2000-231968P	20000917 (60)
US 2000-226279P	20000818 (60)
US 2000-186350P	20000302 (60)
US 2000-184664P	20000224 (60)
US 2000-189874P	20000316 (60)
US 2000-198123P	20000418 (60)
US 2000-227009P	20000823 (60)
US 2000-235484P	20000926 (60)
US 2000-190076P	20000317 (60)
US 2000-209467P	20000607 (60)
US 2000-205515P	20000519 (60)
US 2001-259678P	20010106 (60)

DOCUMENT TYPE: Utility  
 FILE SEGMENT: APPLICATION  
 LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
 ROCKVILLE, MD, 20850  
 NUMBER OF CLAIMS: 24  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 24462

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel ovarian related polynucleotides and the polypeptides encoded by these polynucleotides herein collectively known as "ovarian antigens," and the use of such ovarian antigens for detecting disorders of the ovaries and/or breast, particularly the presence of ovarian and/or breast cancer and ovarian and/or breast cancer metastases. More specifically, isolated ovarian associated nucleic acid molecules are provided encoding novel ovarian associated polypeptides. Novel ovarian polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human ovarian associated polynucleotides and/or polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to the ovaries and/or breast, including ovarian and/or breast cancer, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting the production and function of the polypeptides of the present invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 21 OF 41 USPATEFULL  
 ACCESSION NUMBER: 2000:85170 USPATEFULL  
 TITLE: Neuropeptide like polypeptide zpep17  
 INVENTOR(S): Sheppard, Paul O., Granite Falls, WA, UNITED STATES  
 Bishop, Paul D., Fall City, WA, UNITED STATES

	NUMBER	FIND	DATE
PATENT INFORMATION:	US 2002045210	A1	20020418

APPLICATION INFO.: US 2001-776795 AI 20010205 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-180314P	20000204 (50)
	US 2000-180896P	20000207 (50)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Jennifer K. Johnson, SymoGenetics, Inc, 1201 Eastlake Avenue East, Seattle, WA, 98102	
NUMBER OF CLAIMS:	26	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	12 Drawing Page(s)	
LINE COUNT:	4459	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to polynucleotide and polypeptide molecules for zpep17, a novel secreted protein. The polynucleotides encoding zpep17, may, for example, be used to identify a region of the genome associated with human disease states. The present invention also includes methods for producing the protein, uses therefor and antibodies thereto.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 22 OF 41 USPATFULL

ACCESSION NUMBER: 2002:78729 USPATFULL  
TITLE: Nucleic acids, proteins, and antibodies  
INVENTOR(S): Fosen, Craig A., Laytonsville, MD, UNITED STATES  
Fuben, Steven M., Olney, MD, UNITED STATES  
Barash, Steven C., Rockville, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002042386	AI	20020411
APPLICATION INFO.:	US 2001-764870	AI	20010117 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-179066P	20000131 (50)
	US 2000-180628P	20000204 (50)
	US 2000-214896P	20000628 (50)
	US 2000-217487P	20000711 (50)
	US 2000-225758P	20000814 (50)
	US 2000-220963P	20000726 (50)
	US 2000-217496P	20000711 (50)
	US 2000-225447P	20000814 (50)
	US 2000-218290P	20000714 (50)
	US 2000-225757P	20000814 (50)
	US 2000-226868P	20000822 (50)
	US 2000-216647P	20000707 (50)
	US 2000-225267P	20000814 (50)
	US 2000-216880P	20000707 (50)
	US 2000-225170P	20000814 (50)
	US 2000-221869P	20001208 (50)
	US 2000-225834P	20000907 (50)
	US 2000-224274P	20000911 (50)
	US 2000-224223P	20000911 (50)
	US 2000-227824P	20000830 (50)
	US 2000-224518P	20000814 (50)
	US 2000-226699P	20000909 (50)
	US 2000-224519P	20000814 (50)
	US 2000-220964P	20000726 (50)
	US 2000-224899P	20001020 (50)
	US 2000-229249P	20001117 (50)
	US 2000-226327P	20000929 (50)

US 2000-241785P	20001020 (60)
US 2000-244617P	20001101 (60)
US 2000-235064P	20000814 (60)
US 2000-236368P	20000929 (60)
US 2000-251856P	20001208 (60)
US 2000-251868P	20001208 (60)
US 2000-229344P	20000901 (60)
US 2000-234997P	20000925 (60)
US 2000-229343P	20000901 (60)
US 2000-229345P	20000901 (60)
US 2000-229287P	20000901 (60)
US 2000-229513P	20000901 (60)
US 2000-231413P	20000908 (60)
US 2000-229509P	20000905 (60)
US 2000-236367P	20000929 (60)
US 2000-237039P	20001002 (60)
US 2000-237038P	20001002 (60)
US 2000-236370P	20000929 (60)
US 2000-236803P	20001002 (60)
US 2000-237037P	20001002 (60)
US 2000-237040P	20001002 (60)
US 2000-246960P	20001020 (60)
US 2000-239935P	20001013 (60)

DOCUMENT TYPE: Utility  
 FILE SEGMENT: APPLICATION  
 LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
 ROCKVILLE, MD, 20850  
 NUMBER OF CLAIMS: 24  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 23133

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 23 OF 41 USPATFULL

ACCESSION NUMBER: 2000:78715 USPATFULL

TITLE: Stannocalcin polynucleotides, polypeptides, and methods based thereon

INVENTOR(S): Olsen, Henrik S., Gaithersburg, MD, UNITED STATES  
 Zhang, Ke-Zhou, Brussels, BELGIUM  
 Lindsberg, Perttu, Helsinki, FINLAND  
 Tattisumak, Turgut, Helsinki, FINLAND  
 Kaste, Markku, Vantaa, FINLAND  
 Andersson, Leif C., Helsinki, FINLAND

PATENT ASSIGNEE(S): Human Genome Sciences, Inc., Rockville, MD, UNITED STATES, 20850 (U.S. corporation)

NUMBER	KIND	DATE
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PATENT INFORMATION: US 2002042872 A1 20020411

APPLICATION INFO.: US 2001-940939 A1 20010415 (9)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. WO 2000-US29432, filed

on 26 Oct 2000, UNKNOWN

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-161740P	19991027 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	47	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	12 Drawing Page(s)	
LINE COUNT:	9559	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to human stanniocalcin (STC) polynucleotides, polypeptides, and other Stanniocalcin compositions and to novel methods based thereon. In a specific embodiment, the Stanniocalcin compositions of the invention are used to treat or protect neural cells. Moreover, the present invention relates to vectors, host cells, antibodies, and recombinant and synthetic methods for producing the Stanniocalcin compositions of the invention. Also provided are diagnostic methods for detecting or prognosing diseases, disorders, damage or injury, associated with alterations of the Stanniocalcin compositions of the invention, and to therapeutic methods for treating such diseases, disorders, damage or injury.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 24 OF 41 USPATFULL

ACCESSION NUMBER: 2002:79412 USPATFULL  
TITLE: Nucleic acids, proteins, and antibodies  
INVENTOR(S): Fosen, Craig A., Laytonsville, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES  
Barash, Steven C., Rockville, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002042096	A1	20020411
APPLICATION INFO.:	US 2001-764887	A1	20010117 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-179065P	20000131 (60)
	US 2000-180638P	20000204 (60)
	US 2000-214886P	20000628 (60)
	US 2000-217487P	20000711 (60)
	US 2000-225758P	20000814 (60)
	US 2000-229963P	20000726 (60)
	US 2000-217496P	20000711 (60)
	US 2000-225447P	20000814 (60)
	US 2000-218230P	20000714 (60)
	US 2000-225757P	20000814 (60)
	US 2000-226868P	20000822 (60)
	US 2000-216647P	20000707 (60)
	US 2000-225267P	20000814 (60)
	US 2000-216880P	20000707 (60)
	US 2000-225270P	20000814 (60)
	US 2000-251864P	20001208 (60)
	US 2000-235834P	20000927 (60)
	US 2000-234274P	20000921 (60)
	US 2000-234233P	20000921 (60)
	US 2000-228924P	20000830 (60)
	US 2000-234513P	20000814 (60)
	US 2000-236309P	20000929 (60)
	US 2000-224519P	20000814 (60)

US 2000-220964P	20000726 (60)
US 2000-241899P	20001029 (60)
US 2000-249299P	20001117 (60)
US 2000-236327P	20000909 (60)
US 2000-241785P	20001020 (60)
US 2000-244617P	20001101 (60)
US 2000-225268P	20000814 (60)
US 2000-236368P	20000909 (60)
US 2000-251856P	20001208 (60)
US 2000-251868P	20001208 (60)
US 2000-229344P	20000401 (60)
US 2000-234937P	20000925 (60)
US 2000-229343P	20000901 (60)
US 2000-229345P	20000901 (60)
US 2000-229287P	20000901 (60)
US 2000-229513P	20000905 (60)
US 2000-231413P	20000908 (60)
US 2000-229509P	20000905 (60)
US 2000-236367P	20000909 (60)
US 2000-237039P	20001002 (60)
US 2000-237038P	20001002 (60)
US 2000-236370P	20000909 (60)
US 2000-236802P	20001002 (60)
US 2000-237037P	20001002 (60)
US 2000-237040P	20001002 (60)
US 2000-240960P	20001020 (60)
US 2000-239935P	20001013 (60)

DOCUMENT TYPE: Utility  
 FILE SEGMENT: APPLICATION  
 LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
 ROCKVILLE, MD, 20850  
 NUMBER OF CLAIMS: 24  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 19533  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel liver related polynucleotides and the polypeptides encoded by these polynucleotides herein collectively known as "liver antigens," and the use of such liver antigens for detecting disorders of the liver, particularly the presence of cancer of liver and cancer metastases. More specifically, isolated liver associated nucleic acid molecules are provided encoding novel liver associated polypeptides. Novel liver polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human liver associated polynucleotides and/or polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to the liver, including cancer of liver tissues, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting the production and function of the polypeptides of the present invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 25 OF 41 USPATFULL  
 ACCESSION NUMBER: 2002:56846 USPATFULL  
 TITLE: ABC transport polynucleotides, polypeptides, and antibodies  
 INVENTOR(S): Riben, Steven M., Olney, MD, UNITED STATES  
 Ni, Jian, Germantown, MD, UNITED STATES  
 Moore, Paul A., Germantown, MD, UNITED STATES

NUMBER	KIND	DATE
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PATENT INFORMATION: US 2002037549 A1 20020328  
APPLICATION INFO.: US 2001-757870 A1 20010124 (9)  
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. WO 2000-US19736, filed  
on 20 Jul 2000, UNKNOWN

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-145115P	19990723 (60)
	US 1999-149445P	19990818 (60)
	US 1999-164730P	19991112 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: 12  
EXEMPLARY CLAIM: 1  
LINE COUNT: 12219

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human ABC transport polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human ABC transport polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human ABC transport polypeptides.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 20 OF 41 USPTFLL

ACCESSION NUMBER: 2002:66870 USPTFLL  
TITLE: IL-6-like polynucleotides, polypeptides, and antibodies  
INVENTOR(S): Ruben, Steven M., Olney, MD, UNITED STATES  
Shi, Yanggu, Gaithersburg, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002037543	A1	20020328
APPLICATION INFO.:	US 2001-875016	A1	20010607 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US33134, filed on 7 Dec 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-169838P	19991109 (60)

DOCUMENT TYPE: Utility  
FILE SEGMENT: APPLICATION  
LEGAL REPRESENTATIVE: HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,  
ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: 12  
EXEMPLARY CLAIM: 1  
LINE COUNT: 11587

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human IL-6-like polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human IL-6-like polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human IL-6 like polypeptides.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 27 OF 41 USPTFLL

ACCESSION NUMBER: 2002:48270 USPTFLL

TITLE: Methods for analyzing protein binding events  
INVENTOR(S): Hefti, John J., San Francisco, CA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002028461	A1	20020307
APPLICATION INFO.:	US 2001-923474	A1	20010806 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1999-365580, filed on 2 Aug 1999, GRANTED, Pat. No. US 6287874 Continuation-in-part of Ser. No. US 1999-243144, filed on 1 Feb 1999, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1998-73445P	19980202 (60)
	US 1999-134740P	19990518 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Richard L. Neeley, Clifford B. Perry, Signature BioScience, Inc., 21124 Cabot Boulevard, Hayward, CA, 94545 1130	
NUMBER OF CLAIMS:	45	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	37 Drawing Page(s)	
LINE COUNT:	4041	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a variety of methods of analyzing protein binding events using a system capable of directly detecting protein/ligand complexes based upon the dielectric properties of the complex. The system can be used in a variety of analyses involving protein binding events, such as screening ligand libraries, characterizing protein binding interactions, and identifying ligands. The system can also be utilized in diverse analytical and diagnostic applications.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 28 OF 41 USPTAFULL

ACCESSION NUMBER: 2002:02131 USPTAFULL  
TITLE: 18 Human secreted proteins  
INVENTOR(S): Shi, Yanggu, Gaithersburg, MD, UNITED STATES  
Young, Paul E., Gaithersburg, MD, UNITED STATES  
Ebner, Reinhard, Gaithersburg, MD, UNITED STATES  
Koppert, Daniel R., Centreville, VA, UNITED STATES  
Fukun, Steven M., Olney, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002017466	A1	20020131
APPLICATION INFO.:	US 2001-708826	A1	20010125 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US22350, filed on 15 Aug 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-148759P	19990818 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 2410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	23	
EXEMPLARY CLAIM:	1	
LINE COUNT:	1817	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human secreted proteins and

isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating diseases, disorders, and/or conditions related to these novel human secreted proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 29 OF 41 USPATEFULL

ACCESSION NUMBER: 2002:12261 USPATEULL  
TITLE: Uteroglobin-like polynucleotides, polypeptides, and antibodies  
INVENTOR(S): Ni, Jian, Germantown, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 20020006640	A1	20020117
APPLICATION INFO.:	US 2001 846258	A1	20010502 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US30326, filed on 3 Nov 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-163395P	19991104 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE, ROCKVILLE, MD, 20850	
NUMBER OF CLAIMS:	22	
EXEMPLARY CLAIM:	1	
LINE COUNT:	12076	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human uteroglobin-like polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human uteroglobin-like polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human uteroglobin-like polypeptides.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 30 OF 41 USPATEFULL

ACCESSION NUMBER: 2002:2489 USPATEFULL  
TITLE: Estrogen receptor interacting polynucleotides, polypeptides, and antibodies  
INVENTOR(S): Shi, Yanggu, Gaithersburg, MD, UNITED STATES  
Ruben, Steven M., Olney, MD, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002004489	A1	20020110
APPLICATION INFO.:	US 2001-788660	A1	20010221 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000 US22351, filed on 15 Aug 2000, UNKNOWN		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-143717P	1999-16 (60)
	US 2000-144026P	20000314 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	HUMAN GENOME SCIENCES INC, 9410 KEY WEST AVENUE,	

ROCKVILLE, MD, 20850

NUMBER OF CLAIMS: 1  
EXEMPLARY CLAIM: 1  
LINE COUNT: 11257

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to novel human RIP polypeptides and isolated nucleic acids containing the coding regions of the genes encoding such polypeptides. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human RIP polypeptides. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human RIP polypeptides.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 31 OF 41 USPATFULL

ACCESSION NUMBER: 2002:75196 USPATFULL  
TITLE: Bio assay device and test system for detecting molecular binding events  
INVENTOR(S): Hefti, John, San Francisco, CA, United States  
PATENT ASSIGNEE(S): Signature BioScience, Inc., Hayward, CA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6368795	B1	20020409
APPLICATION INFO.:	US 1999-243194		19990201 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 1998-73445P	19980202 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Chin, Christopher L.	
LEGAL REPRESENTATIVE:	Neeley, Richard L., Perry, Clifford B.	
NUMBER OF CLAIMS:	13	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	51 Drawing Figure(s); 28 Drawing Page(s)	
LINE COUNT:	3253	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Systems and methods for detecting molecular binding events and other environmental effects using the unique dielectric properties of the bound molecular structure or structures are presented. A molecular binding layer is coupled along the surface of a signal path. A test signal is propagated along the signal path, whereby the test signal couples to the molecular binding layer, and in response, exhibits a signal response.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 32 OF 41 USPATFULL

ACCESSION NUMBER: 2002:9766 USPATFULL  
TITLE: Method and apparatus for detecting molecular binding events  
INVENTOR(S): Hefti, John, San Francisco, CA, United States  
PATENT ASSIGNEE(S): Signature BioScience, Inc., Hayward, CA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6368795	B1	20020115
APPLICATION INFO.:	US 1999-365578		19990802 (9)
RELATED APPLN. INFO.:	Continuation-in part of Ser. No. US 1999-243194, filed on 1 Feb. 1999		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1998-73445P	19980202 (50)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Chin, Christopher L.	
LEGAL REPRESENTATIVE:	Perry, Clifford B., Neeley, Richard L.	
NUMBER OF CLAIMS:	20	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	51 Drawing Figure(s); 28 Drawing Page(s)	
LINE COUNT:	3281	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Systems and methods are presented for detecting molecular binding events and other environmental effects using the unique dielectric properties of the bound molecular structure or structures. A molecular binding region is coupled along the surface of a signal path. A test signal is propagated along the signal path, whereby the test signal couples to the molecular binding region, and in response, exhibits a signal response.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 33 OF 41 EUROPATFULL COPYRIGHT 2002 WILA

GRANTED PATENT - ERTEILTES PATENT - BREVET DELIVRE

ACCESSION NUMBER:	786087	EUROPATFULL	EW 200209	FS PS																														
TITLE:	POLYPEPTIDE-DENDRIMER COMPLEXES. POLYPEPTID-DENDRIMER KOMPLEXE. COMPLEXES DE POLYPEPTIDES-DENDRIMES.																																	
INVENTOR(S):	SINGH, Pratap, 19111 S.W. 69th Street, Miami, FL 33193, US																																	
PATENT ASSIGNEE(S):	DADE BEHRING INC., 1717 Deerfield Road, Deerfield, Illinois 60015, US																																	
PATENT ASSIGNEE NO:	1905941																																	
AGENT:	Helbing, Joerg, Dr. Dipl.-Chem. et al., Patentanwaelte von Kreisler-Selting-Werner, Postfach 10 22 41, 50462 Koeln, DE																																	
AGENT NUMBER:	80653																																	
OTHER SOURCE:	BEP2002015 EP 0786087 B1 0029																																	
SOURCE:	Wila EPS-2002-H02-T2																																	
DOCUMENT TYPE:	Patent																																	
LANGUAGE:	Anmeldung in Englisch; Veroeffentlichung in Englisch																																	
DESIGNATED STATES:	F AT; F BE; F CH; F DE; F DK; F ES; F FI; F FR; F GB; R GR; F IE; F IT; F LI; F LU; F MC; F NL; F PT; R SE																																	
PATENT INFO.PUB.TYPE:	EPRI EUROPAEISCHE PATENTSCHRIEFT (Internationale Anmeldung)																																	
PATENT INFORMATION:	<table border="0"> <thead> <tr> <th>PATENT NO</th> <th>KIND</th> <th>DATE</th> </tr> </thead> <tbody> <tr> <td>EP 786087</td> <td>B1</td> <td>20020227</td> </tr> <tr> <td></td> <td></td> <td>19970730</td> </tr> <tr> <td>APPLICATION INFO.:</td> <td>EP 1996-927393</td> <td>19960804</td> </tr> <tr> <td>PRIORITY APPLN. INFO.:</td> <td>US 1996-514075</td> <td>19960811</td> </tr> <tr> <td>RELATED DOC. INFO.:</td> <td>WO 96-0313957</td> <td>960809 INTAKZ</td> </tr> <tr> <td></td> <td>WO 9707498</td> <td>970227 INTENT</td> </tr> <tr> <td>REFERENCE PAT. INFO.:</td> <td>WO 89 01174 A</td> <td>WO 94-18693 A</td> </tr> <tr> <td></td> <td>WO 91 27902 A</td> <td>WO 95 18641 A</td> </tr> <tr> <td>REF. NON-PATENT LIT.:</td> <td colspan="2">BIOCONJUGATE CHEMISTRY, vol. 1, no. 5, 1 September 1990, pages 305-208, XP00017464 ROBERTS J C ET AL: "USING STARBURST DENDRIMERS AS LINKER MOLECULES TO RADIOLEBEL ANTIBODIES" ABSTRACTS OF PAPERS AMERICAN CHEMICAL SOCIETY, vol. 211, no. 12, 14 - 28 March 1996, NEW ORLEANS, page B1CT 193 XP00242032 P. SINGH: "Coupling of multiple proteins to starburst dendrimers."</td> </tr> </tbody> </table>				PATENT NO	KIND	DATE	EP 786087	B1	20020227			19970730	APPLICATION INFO.:	EP 1996-927393	19960804	PRIORITY APPLN. INFO.:	US 1996-514075	19960811	RELATED DOC. INFO.:	WO 96-0313957	960809 INTAKZ		WO 9707498	970227 INTENT	REFERENCE PAT. INFO.:	WO 89 01174 A	WO 94-18693 A		WO 91 27902 A	WO 95 18641 A	REF. NON-PATENT LIT.:	BIOCONJUGATE CHEMISTRY, vol. 1, no. 5, 1 September 1990, pages 305-208, XP00017464 ROBERTS J C ET AL: "USING STARBURST DENDRIMERS AS LINKER MOLECULES TO RADIOLEBEL ANTIBODIES" ABSTRACTS OF PAPERS AMERICAN CHEMICAL SOCIETY, vol. 211, no. 12, 14 - 28 March 1996, NEW ORLEANS, page B1CT 193 XP00242032 P. SINGH: "Coupling of multiple proteins to starburst dendrimers."	
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L33 ANSWER 34 OF 41 PCTFULL COPYRIGHT 2002 Univentio  
 ACCESSION NUMBER: 2002036624 PCTFULL ED 20020523 EW 200219  
 TITLE (ENGLISH): METHODS AND COMPOSITIONS RELATING TO FORTILIN, AN  
 ANTI-APOPTOTIC MOLECULE, AND MODULATORS OF FORTILIN  
 TITLE (FRENCH): PROCEDES ET COMPOSITIONS ASSOCIES A LA FORTILINE, UNE  
 MOLECULE ANTI-APOPTOTIQUE, ET MODULATEURS DE FORTILINE  
 INVENTOR(S): FUJISE, Kenichi; YEH, Edward  
 PATENT ASSIGNEE(S): BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM, for  
 all designates States except US; FUJISE, Kenichi, for  
 US only; YEH, Edward, for US only  
 AGENT: SHISHIMA, Gina, N.  
 LANGUAGE OF PUBL.: English  
 LANGUAGE OF FILING: English  
 DOCUMENT TYPE: Patent  
 PATENT INFORMATION:

NUMBER	KIND	DATE
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DESIGNATED STATES:	WO 2002036624	A2 20020510
	AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR	
	CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HF HU ID	
	IL IN IS JP KE KG KP KR KZ LC LK LP LS LT LU LV MA MD	
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	SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW GH GM KE LS	
	MW ME SD SL SZ TZ UG ZW AM AZ BY BG BZ MD RU TJ TM AT	
	BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR	
	BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG	

APPLICATION INFO.: WO 2001/US42985 A 20011030  
 PRIORITY INFO.: US 2000 60/244,416 20001030

ABEN The polypeptide Fortilin (also known as Translationally Controlled  
 Tumour Protein, TCTP) specifically interacts with p53, a tumor  
 suppressor involved in the induction of apoptosis and the normal growth  
 regulation of a cell. Fortilin also specifically binds MCL1 (Myeloid  
 Cell Leukemia 1). Fortilin has the ability to prevent apoptosis, which  
 may be unregulated in hyperproliferative cells. The present invention is  
 directed at compositions and methods involving a Fortilin modulator,  
 which can induce apoptosis, for the prevention, treatment, or diagnosis  
 of hyperproliferative diseases and conditions, including cancer and  
 atherosclerosis. It is directed also at compositions and methods  
 involving Fortilin, which can inhibit apoptosis, for the treatment of  
 diseases and condition characterized by apoptosis, including certain  
 vascular conditions.

ABFR Le polypeptide fortiline (egalement appele proteine tumorale de  
 regulation de traduction, TCTP) interagit specifiquement avec p53, un  
 supprimeur de tumeur intervenant dans l'induction de l'apoptose et la  
 regulation de la croissance normale d'une cellule. La fortiline se lie  
 aussi specifiquement a MCL1 (leucemie myeloide 1). La fortiline est  
 capable de prevenir l'apoptose, qui peut etre deregulee dans des cellules  
 hyperproliferatives. L'invention concerne des compositions et des  
 procedes comprenant un modulateur de fortiline, capable d'induire  
 l'apoptose, pour prevenir, traiter ou diagnostiquer des maladies ou des  
 affections hyperproliferatives, y compris le cancer et l'atherosclerose  
 ; ainsi que des compositions et des procedes comprenant la fortiline,  
 capable d'inhiber l'apoptose, pour traiter des maladies et affections  
 caracterisees par l'apoptose, y compris certaines affections  
 vasculaires.

L33 ANSWER 35 OF 41 USPATFULL  
 ACCESSION NUMBER: 2001:15781 USPATFULL  
 TITLE: Methods for analyzing protein binding events  
 INVENTOR(S): Hoffi, John, San Francisco, CA, United States  
 PATENT ASSIGNEE(S): Signature BioScience, Inc., Hayward, CA, United States  
 (U.S. corporation)

NUMBER	KIND	DATE
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PATENT INFORMATION: US 6287874 B1 20010911  
APPLICATION INFO.: US 1999-365580 19990802 (9)  
RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1999-243194, filed  
on 1 Feb 1999

	NUMBER	DATE
PRIORITY INFORMATION:	US 1998-73445P	19980202 (60)
	US 1999-134740P	19990518 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	GRANTED	
PRIMARY EXAMINER:	Horlick, Kenneth R.	
ASSISTANT EXAMINER:	Strzelecka, Teresa	
LEGAL REPRESENTATIVE:	Ausenhuis, Scott L., Perry, Clifford B., Neeley, Richard L.	
NUMBER OF CLAIMS:	45	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	44 Drawing Figure(s); 33 Drawing Page(s)	
LINE COUNT:	4099	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a variety of methods of analyzing protein binding events using a system capable of directly detecting protein/ligand complexes based upon the dielectric properties of the complex. The system can be used in a variety of analyses involving protein binding events, such as screening ligand libraries, characterizing protein binding interactions, and identifying ligands. The system can also be utilized in diverse analytical and diagnostic applications.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 36 OF 41 USPATFULL

ACCESSION NUMBER: 2000:87731 USPATFULL  
TITLE: Methods and compositions for using membrane-penetrating proteins to carry materials across cell membranes  
INVENTOR(S): Draper, Rockford, Plano, TX, United States  
PATENT ASSIGNEE(S): Board of Regents, The University of Texas Systems, Austin, TX, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6086900		20000711
APPLICATION INFO.:	US 1998-47148		19980324 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 1997-42056P	19970326 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Guze, David	
LEGAL REPRESENTATIVE:	Arnold, White & Durkee	
NUMBER OF CLAIMS:	62	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	8 Drawing Figure(s); 6 Drawing Page(s)	
LINE COUNT:	2729	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides methods and compositions delivery of agents into the cytoplasm of cells. Particularly, it concerns the use of membrane penetrating toxin proteins to deliver drugs to the cytoplasm of target cells.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 37 OF 41 USPATFULL

ACCESSION NUMBER: 2000:84049 USPATFULL

TITLE: Polypeptide: dendrimer complexes  
 INVENTOR(S): Singh, Pratap, Wilmington, DE, United States  
 Lin, Spencer, Granger, IN, United States  
 Moll, III, Fred, Pembroke Pines, FL, United States  
 PATENT ASSIGNEE(S): Dade Behring Inc., Deerfield, IL, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6082708		20000704
APPLICATION INFO.:	US 1995-514075		19950811 (8)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Wortman, Donna C.		
LEGAL REPRESENTATIVE:	Lundquist, Ronald C, Tymeson, Cynthia G		
NUMBER OF CLAIMS:	38		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	5 Drawing Figure(s); 1 Drawing Page(s)		
LINE COUNT:	1674		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Compositions are disclosed, comprising dendrimers to which a first polypeptide is controllably coupled. Such polypeptide-dendrimer compositions are effective for controllably coupling a second polypeptide to the dendrimer. The first and second polypeptides have separate and distinct defined biological activities, for example, two antibodies with first and second binding specificities or an antibody and an enzymatic label. Such compositions are useful as indicators in specific binding assays, e.g., immunoassays. Methods for sequentially coupling two different polypeptides to a dendrimer to form compositions of the invention also are disclosed.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 38 OF 41 MEDLINE  
 ACCESSION NUMBER: 90062164 MEDLINE  
 DOCUMENT NUMBER: 90062164 PubMed ID: 2584219  
 TITLE: The reactivity of **sulphydryl groups** of bovine cardiac **troponin C**.  
 AUTHOR: Fuchs F; Liou Y M; Grabarek Z  
 CORPORATE SOURCE: Department of Physiology, University of Pittsburgh School of Medicine, Pennsylvania 15261.  
 CONTRACT NUMBER: AR-19551 (NIAMS)  
 SOURCE: E-37-HL 05949 (NHLBI)  
 JOURNAL: JOURNAL OF BIOLOGICAL CHEMISTRY, (1989 Dec 5) 264 (34) 20344-9.  
 PUB. COUNTRY: United States  
 LANGUAGE: English  
 FILE SEGMENT: Priority Journals  
 ENTRY MONTH: 199001  
 ENTRY DATE: Entered STN: 19900373  
 Last Updated on STN: 19970103  
 Entered Medline: 19900108

AB bovine cardiac **troponin C** (cTnC) contains 2 cysteine residues, Cys-35 located in the nonfunctional Ca<sup>2+</sup>-binding loop I and Cys-84 in the N-terminal segment of the central helix. We have studied the reactivity of Cys residues in cTnC with 5,5'-dithiobis(2-nitrobenzoic acid) (DTNB) and 7-diethylamino-2-(4'-maleimidylphenyl)-4-methylcoumarin (CPM). The latter compound fluoresces only when reacted with the protein. The reaction with DTNB followed second order kinetics with respect to DTNB, the rate constants being 3.27 s<sup>-1</sup> M<sup>-1</sup> and 1.82 s<sup>-1</sup> M<sup>-1</sup> in the presence and absence of Ca<sup>2+</sup>, respectively. These rates are much slower than the rate of reaction with Cys-23 of skeletal TnC (sTnC) or with the urea-denatured cTnC, indicating that both Cys residues are partly buried within the



structure of the protein. The increase in reactivity was induced by binding of  $\text{Ca}^{2+}$  to the single low affinity  $\text{Ca}^{2+}$  binding site (site II). The fluorescence increase upon reaction of cTnC with CPM in the absence of  $\text{Ca}^{2+}$  could be fitted with a single exponential equation indicating that both cysteine residues are equally available to the reagent. The reaction in the presence of  $\text{Ca}^{2+}$  was biphasic. Analysis of CNBr fragments of cTnC labeled with CPM under various conditions indicated that in the presence of  $\text{Ca}^{2+}$  the reactivity of Cys-84 is increased while that of Cys-35 is slightly decreased. This finding is consistent with the model of Herzberg et al. (Herzberg, O., Moulton, J., and James, M. N. G. (1986) J. Biol. Chem. 261, 2638-2644) and the data of Ingraham and Hodges (Ingraham, R. H., and Hodges, R. S. (1988) Biochemistry 27, 5891-5898), suggesting that the  $\text{Ca}^{2+}$ -induced conformational change in the N-terminal half of TnC involves separation of the helix C from the central helix, thereby increasing the accessibility of Cys-84. The slow overall kinetics, however, indicates that the structure in the vicinity of Cys residues is relatively compact regardless of  $\text{Ca}^{2+}$ . We interpret the increase in reactivity towards CPM as consistent with a  $\text{Ca}^{2+}$ -induced exposure of a hydrophobic pocket in the vicinity of Cys-84.

L33 ANSWER 39 OF 41 USPATFULL

ACCESSION NUMBER: 86:34211 USPATFULL  
 TITLE: Protein kinase enzyme AUT-PK 500 and a radioimmunoassay for detection of neoplasia  
 INVENTOR(S): Sharma, Rameshwar K., Memphis, TN, United States  
 PATENT ASSIGNEE(S): The University of Tennessee Research Corp., Knoxville, TN, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4594319		19860610
APPLICATION INFO.:	US 1984-590712		19840319 (6)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Wiseman, Thomas G.		
ASSISTANT EXAMINER:	Moskowitz, M.		
LEGAL REPRESENTATIVE:	Neuner, George W., Linek, Ernest V.		
NUMBER OF CLAIMS:	27		
EXEMPLARY CLAIM:	1,6		
NUMBER OF DRAWINGS:	10 Drawing Figure(s); 7 Drawing Page(s)		
LINE COUNT:	1019		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention is directed to AUT-PK 500, a novel autophosphorylating protein kinase, to the **purification** and characterization of AUT-PK 500 from rat adrenocortical carcinoma, to the use of AUT-PK 500 as a marker for neoplasia cells, and to a radioimmunoassay for detecting AUT PK 500 in neoplasia cells.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L33 ANSWER 40 OF 41 MEDLINE

ACCESSION NUMBER: 83135739 MEDLINE  
 DOCUMENT NUMBER: 83135739 PubMed ID: 6826548  
 TITLE: Hydrodynamic properties of bovine cardiac **troponin** -I and **troponin**-T.  
 AUTHOR: Byers D M; Kay C M  
 SOURCE: JOURNAL OF BIOLOGICAL CHEMISTRY, (1983 Mar 10) 258 (5) 2351-4.  
 Journal Code: 2385121R. ISSN: 0021-9258.  
 PUB. COUNTRY: United States  
 Journal; Article; (JOURNAL ARTICLE)  
 LANGUAGE: English  
 FILE SEGMENT: Priority Journals  
 ENTRY MONTH: 198304  
 ENTRY DATE: Entered STN: 19900318

Last Updated on STN: 19990129

Entered Medline: 19830407

AB Bovine cardiac **troponin**-I (TN-I) and **troponin**-T (TN-T) have been examined in solution using ultracentrifugation, gel filtration, and viscosity. A new method of purifying TN-T, employing hydroxylapatite chromatography in 6 M urea, is reported. Cardiac TN-T (Mr = 36,000) undergoes a reversible, concentration-dependent association in nondenaturing buffers, probably to a tetramer. The Stokes radius (Rs) of aggregated TN-T, determined by sedimentation velocity and gel chromatography on Sephacryl S-300, is 80 A and the reduced viscosity of the subunit ranges from 20 to 25 ml/g at protein concentrations between 0.5 and 2.5 mg/ml. These data suggest that TN-T forms highly asymmetric aggregates in solution. Bovine cardiac TN-I also has a tendency toward self-association, but is essentially monomeric (Mr = 23,000) at protein concentrations below 1 mg/ml. The presence of reducing agent is necessary to avoid intermolecular disulfide bond formation. From gel filtration experiments, the value of Rs is 29 A indicating that TN-I is a moderately asymmetric protein (frictional ratio = 1.5). Similar properties are observed when both **sulfhydryl groups** of TN-I are modified by carboxamidomethylation.

L33 ANSWER 41 OF 41 MEDLINE  
ACCESSION NUMBER: 71011997 MEDLINE  
DOCUMENT NUMBER: 71011997 PubMed ID: 4248628  
TITLE: A study of the role of **sulfhydryl groups**  
in the interaction of **troponin** and myofibrils.  
AUTHOR: Parker C J Jr; Kilbert L H Jr  
SOURCE: ARCHIVES OF BIOCHEMISTRY AND BIOPHYSICS, (1970 Oct) 140 (2)  
326-33.  
Journal code: 0372430. ISSN: 0003-9861.  
PUB. COUNTRY: United States  
Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 197012  
ENTRY DATE: Entered STN: 19900101  
Last Updated on STN: 19900101  
Entered Medline: 19701209

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1. *Journal of the American Medical Association*, 2000; 283: 2689-2694.

FAMILY & INT. NUM. TROPNIN: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002004512	A2	20020117	WO 2001-US21817	20010710
WO 2002004512	A3	20020516		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, EC, EE, ES, FI, FR, GB, GR, HU, IL, IN, JP, KE, KG, KH, KR, KZ, LA, LB, LC, LI, LU, LV, MA, MG, MK, MN, MU, MV, MW, MY, NZ, OM, PA, PE, PG, PH, PK, PL, PT, RO, RU, SD, SE, SI, SK, SL, SM, SN, SR, ST, SV, SW, SY, TD, TH, TJ, TK, TL, TR, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, FG, FZ, MD, FH, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, NI, SL, SS, TG, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, ME, NE, SN, TD, TG

US 2002064835 A1 20020130 US 2001-904398 20010710  
 US 2002055147 A1 20020509 US 2001-994619 20011130

PRIORITY APPLN. INFO.:  
 US 2000-210669P P 20000311  
 US 2001-904398 A1 20010710

AB The invention is directed to methods for purifying **troponin I**, particularly recombinant **troponin I** produced in a bacterial expression system. Recombinant **troponin I** can be advantageously purified after reversibly protecting the free sulfhydryl groups, e.g. by forming sulfates. In a specific example, **troponin I** reacted with sodium tetrathionate yielded sulfatolyzed **troponin I**, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the sulfhydryl groups yields a highly purified product ready for refolding.

L67 ANSWER 2 OF 6 USPATEFULL

ACCESSION NUMBER: 2002:126323 USPATEFULL  
 TITLE: Purification of human **troponin I**

INVENTOR(S):  
 Conn, Gregory, Cary, NJ, UNITED STATES  
 Keardson, Brian, Seattle, WA, UNITED STATES  
 Beng, Xiantang, Northborough, MA, UNITED STATES  
 Chang, Chenming, Blacksburg, VA, UNITED STATES

PATENT ASSIGNEE(S):  
 Diosynth RTP, Inc. (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2001-904398	A1	20020030
APPLICATION INFO:	US 2001-123345	A1	20010110

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-210669P	20000710 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	DARBY & DARBY LLP, 300 Third Avenue, New York, NY, 10017	
NUMBER OF CLAIMS:	1	
EXEMPTARY CLAIM:	1	
NUMBER OF DRAWINGS:	11 drawing labels	
LINK COUNT:	566	

TAG INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention is directed to methods for purifying **Troponin I**, particularly recombinant **Troponin I** produced in a bacterial expression system. Recombinant **Troponin I** can be advantageously purified after reversibly protecting the free sulfhydryl groups, e.g. by forming sulfates. In a specific example, **Troponin I** reacted with sodium tetrathionate yielded sulfatolyzed **Troponin I**, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the sulfhydryl groups yields a highly purified product ready for refolding.

AS INDEXING IS AVAILABLE FOR THIS PATENT.

LET ANSWER 5 OF 6 DISAPPEAR!

ACCESSION NUMBER: 20711694 TRIATF001  
TITLE: Purification of human troponin

INVENTOR(S): Deng, Gregory, Cary, NC, UNITED STATES  
Fearndon, Brian, Seattle, WA, UNITED STATES  
Deng, Xianfang, Northborough, MA, UNITED STATES  
Fhang, Chenming, Blacksburg, VA, UNITED STATES  
PATENT ASSIGNEE(S): Licsynth RTP, Inc. (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002051145	A1	20020509
APPLICATION INFO.:	US 2001-998619	A1	20011130 (9)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 2001-903398, filed on 10 Jul 2001, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	NY 1000-217000	20 JUL 1967
DOCUMENT TYPE:	Priority	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	DARBY & DARBY P.C., 805 Third Avenue, New York, NY, 10022	

NUMBER OF CLAIMS: 20  
EXEMPLARY CLAIM: 1  
NUMBER OF DRAWINGS: 11 Drawing Page(s)  
LINE COUNT: 260  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention is directed to methods for purifying Troponin I, particularly recombinant Troponin I produced in a bacterial expression system. Recombinant Troponin I can be advantageously purified after reversibly protecting the free sulfhydryl groups, e.g., by forming sulfates. In a specific example, Troponin I reacted with sodium tetrathionate yielded sulfato-lyzed Troponin I, which was purified by chromatography or ion-exchange, followed by hydrophobic interaction chromatography. Finally, deprotection of the sulfhydryl groups yields a highly purified product ready for refolding.

NO INDEXING IS AVAILABLE FOR THIS PATENT.

L67 ANSWER 4 OF 6 WPILS (C) 2002 THOMSON DERWENT

ACCESSION NUMBER: 25 A-154921 1201 WPIDS  
 ANAL. NO. CIA: 75 A-4849  
 TITLE: Purified troponin I separates  
 subunit and troponin I is

and chromatography on a thin exchanger after previously protecting the free sulphydryl groups.

PERKENT CLASS: B14 D16  
INVENTOR(S): COHN, S; PEARDON, P; ZENG, X; ZHANG, C  
PATENT ASSIGNEE(S): (DIOS-N) DIOSYNTH FTP INC  
PRIORITY INVENT: 05  
PRIORITY INVENTOR: N;  
PRIORITY INVENTOR: N;

Table 1. <i>Continued</i>	Reference	Study design	Study period	Study location	Study population	Sample size	Outcome
10	10	Case-control	1995-1996	USA	Healthcare workers	100	100%
11	11	Case-control	1995-1996	USA	Healthcare workers	100	100%
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58	58	Case-control	1995-1996	USA	Healthcare workers	100	100%
59	59	Case-control	1995-1996	USA	Healthcare workers	100	100%

7. 1. 1971

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GG GH GI GJ GK GL GM GN GP GR GS GT GU GV GW GY HA HB HC HD HE HF HG HH HI  
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IJ IK IL IM IN IO IP IQ IR IS IT IU IV IW IX IY JY JA JB JC JD JE JF JG JH JI  
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KJ KK KL KM KN KO KP KQ KR KS KT KU KV KW KY KZ LY LA LB LC LD LE LF LG LH LI  
LJ LK LL LM LN LO LP LQ LR LS LT LU LV LW LY LZ MY MA MB MC MD ME MF MG MH MI  
MJ MK ML MN MO MP MQ MR MS MT MU MV MW MY MZ NY NA NB NC ND NE NF NG NH NI  
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QJ QK QL QM QN QO QP QQ QR QS QT QU QV QW QY QZ RY RA RB RC RD RE RF RG RH RI  
RJ RK RL RM RN RO RP RQ RR RS RT RU RV RW RY RZ SY SA SB SC SD SE SF SG SH SI  
SJ SK SL SM SN SO SP SQ SR SS ST SU SV SW SY SZ TY TA TB TC TD TE TF TG TH TI  
TJ TK TL TM TN TO TP TQ TR TS TT TU TV TW TY TZ UY UA UB UC UD UE UF UG UH UI  
UJ UK UL UM UN UO UP UQ UR US UT UV UW UY UZ VY VA VB VC VD VE VF VG VH VI  
VJ VK VL VM VN VO VP VQ VR VS VT VU VW VY VZ WY WA WB WC WD WE WF WG WH WI  
WJ WK WL WM WN WO WP WQ WR WS WT WU WV WY WZ XY XA XB XC XD XE XF XG XH XI  
XJ XK XL XM XN XO XP XQ XR XS XT XU XV XW XY XZ YY YA YB YC YD YE YF YG YH  
YI YJ YK YL YM YN YO YP YQ YR YS YT YU YV YW YY YZ ZY ZA ZB ZC ZD ZE ZF ZG  
ZH ZI ZJ ZK ZL ZM ZN ZO ZP ZQ ZR ZS ZT ZU ZV ZW ZY ZZ

Figure 1. The effect of the concentration of the *Agrobacterium* suspension on the transformation efficiency of *Agrobacterium* strains. The number of transformed cells was determined by the number of colonies growing on the selective medium. The results are the mean of three independent experiments. Error bars represent the standard deviation.

## APPLICATION DETAILS:

PATENT NO	KIND	APPLICATION	DATE
WO 2002/045112 A		WO 2001-998619	20010710
AT 2001-0334 A		AT 2001-0334 A	20010710
US 2002/055141 A1	Provisional	US 2001-998619	20010710
	Cont. of	US 2001-998619	20011130

## FILING DETAILS:

PATENT NO	KIND	PATENT NO
AU 2001/0334 A	Based on	WO 2001/0334 A

PRIORITY APPLN. INFO: US 2000-217069P 20000710; US 2001-903398  
20010710; US 2001-998619 20011130

AN 2002-154921 [23] WPIDS

AB WO 200204512 A UPAB: 20020402

NOVELTY - Preparing **troponin I**, comprising protecting free sulphydryl groups of **troponin I** under reducing conditions, and **troponin I** is then purified by subjecting **troponin I** comprising sulphydryl protecting groups to chromatography, is new.

DETAILED DESCRIPTION - An INDEPENDENT CLAIM is also included for **troponin I** comprising sulphydryl protecting groups.

ACTIVITY - Cytostatic.

MECHANISM OF ACTION - Inhibitor of angiogenesis. No supporting data is given.

USE - The method is useful for purifying **troponin I**, particularly recombinant **troponin I**. The highly purified **troponin I**, preferably in a reduced state, is useful for antibody generation, as a control or standard immunoassay reagent, or to inhibit angiogenesis important in treating various cancers.

ADVANTAGE - Protection of sulphydryl groups during **troponin I** preparation eliminates the costly need for maintaining non-reducing conditions throughout protein preparation, **purification** and storage, and need for reducing agents. The **sulphydryl-protected** troponin does not form intrachain or interchain disulfide crosslinks. Overall yield of **troponin I** from the multi-step **purification** was greater than 5% at purity levels of greater than 95%. Deprotection of the sulphydryl groups yields a highly purified product ready for refolding.

Dwg. 011

100 ANSWER 1 OF 6 IFIPAT COPYRIGHT 2002 IFI

AN 2002/055141 IFIPAT:IFIPAT:IFIPAT

TITLE: PURIFICATION OF HUMAN TROPONIN

I

INVENTOR(S):

Wang; Gregory, Cary, US, US

Reardon; Brian, Seattle, WA, US

Zeng; Xianfang, Northborough, MA, US

Zhang; Chenming, Blacksburg, VA, US

PATENT ASSIGNEE(S):

AGENT: DIO SYNTH RTP, INC.

HARRY & DARRY P.C., 205 Third Avenue, New York, NY,

1001, US

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PATENT INFORMATION: US 2002064835 A1 20020130

APPLICATION INFORMATION: US 2001-903398 20010710

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NUMBER DATE

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FAMILY OR PRIORITY NO.  
DOCUMENT TYPE:

FILE SEGMENT:

NUMBER OF CLAIMS:

US 2000-0110,970 (Provisional)

US 2000-0110,970 (Provisional)

INVENTOR:

Patent Application - First Publication

CHEMICAL

APPLICATION

20 11 Figure(s).

#### DESCRIPTION OF FIGURES:

FIGS. 1A and 1B. A. Proposed reaction for oxidative sulfitolysis. B. Cleavage of disulfide bond by sodium sulfite to form the S-sulfite derivative.

FIG. 2. Preparation and washing of TnI-containing inclusion bodies.

FIG. 3. Summary of rTroponin-I preparation.

FIG. 4. Q-Sepharose FF chromatography of **Troponin I**. Buffer

A: 6 M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6 M urea, 25 mM Tris-HCl, pH 7.5, 2 M NaCl; Gradient: Step, 0% B for the flow-through and 100% B for the strip; and Flow rate: 150 ml/min.

FIG. 5. 30' ml Q-Sepharose FF chromatography. Buffer A: 6 M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6 M urea, 25 mM Tris-HCl, pH 7.5, 2 M NaCl; Gradient: Step, 4% B for elution and 6% B for strip; and Flow rate: 20 ml/min.

FIG. 6. SDS-PAGE analysis troponin lot after anion exchange steps no. 1 and no. 2. 16% tris-glycine gel, under non-reducing conditions. A-H refer to lanes in the SDS-PAGE gel. A. Sulfitolyzed troponin Lot 3L4 standard; B. solubilized inclusion bodies; C. sulfitolyzed inclusion bodies (AEX No. 1 load); D. anion exchange no. 1 flowthrough; E. anion exchange no. 1 salt eluate; F. anion exchange no. 2 load; G. anion exchange no. 2 flowthrough; and, H. anion exchange no. 2 100 mM NaCl eluate.

FIG. 7. Toyopearl 650 M (phenyl) HIC chromatograph. Buffer A: 6 M urea, 25 mM Tris-HCl, pH 7.5, 1 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>; Buffer B: 6 M urea, 25 mM Tris-HCl, pH 7.5; Gradient: Step, 100% B for the flow-through and 0% B for strip; and Flow rate: 10 ml/min.

FIGS. 8. SDS-PAGE analysis troponin lot after hydrophobic interaction chromatography in 16% tris-glycine gel, under non-reducing conditions. A-F refers to lanes in the SDS-PAGE gel. A. Sulfitolyzed troponin Lot 3L4 standard; B. AEX step no. 2, troponin eluate pool; C. HIC load (w/1M ammonium sulfate); D. HIC flowthrough (troponin product); E. HIC low salt eluate (column strip); F. lot 3L4 sulfitolyzed troponin product.

FIG. 9. Quantitation of rTnI on column.

FIG. 10. **Troponin I** LysC mapping.

FIG. 11. SDS-PAGE analysis of sulfitolyzed troponin reduction with dithiothreitol for 45 mins. at ambient temperature. One mg/ml TnI in 6 M urea, 25 mM tris, 0.15 M NaCl pH 7.5, run on 16% tris-glycine gel. 1. 10%, Mark 12 MW Plus; 2. 9%, sulfitolyzed TnI; 3. 0.05 mM DTT; 4. 0.10 mM DTT; 5. 0.2 mM DTT; 6. 0.5 mM DTT; 7. 1.0 mM DTT; 8. 1.5 mM DTT.

AB The invention is directed to methods for purifying **Troponin I**, particularly recombinant troponin I produced in a bacterial expression system. Recombinant troponin I can be purified by various methods after reversibly protecting the free sulfhydryl groups, e.g., by forming sulfates. In a specific example, Troponin I reacted with sodium tetrafluoroborate yielded sulfitolyzed Troponin I, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the sulfhydryl groups yields a highly purified product ready for refolding.

FIGS. 1A and 1B.

FIGS. 1A and 1B. A. Proposed reaction for oxidative sulfitolysis. B. Cleavage of disulfide bond by sodium sulfite to form the S-sulfite derivative.

FIG. 2. Preparation and washing of TnI-containing inclusion bodies.

FIG. 3. Summary of rTroponin-I preparation.

FIG. 4. Q-Sepharose FF chromatography of **Troponin I**.

Buffer A: 6 M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6 M urea, 25 mM Tris-HCl, pH 7.5, 2 M NaCl; Gradient: Step, 0% B for the flow-through and 100% B for the strip; and Flow rate: 150 ml/min.

FIG. 5. 30' ml Q-Sepharose FF chromatography. Buffer A: 6 M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6 M urea, 25 mM Tris-HCl, pH 7.5, 2 M NaCl; Gradient: Step, 4% B for elution and 6% B for strip; and Flow rate: 20 ml/min.

FIG. 6. SDS-PAGE analysis troponin lot after anion exchange steps no. 1 and no. 2 in 16 $\times$  tris-glycine gel, under nonreducing conditions. A-H refer to lanes in the SDS-PAGE gel. A. Sulfitolyzed troponin Lot 3L4 standard; B. solubilized inclusion bodies; C. sulfitolyzed inclusion bodies; A+E No. 1 eluate; D. anion exchange no. 1 flowthrough; E. anion exchange no. 1 salt eluate; F. anion exchange no. 2 flowthrough; G. anion exchange no. 2 salt eluate; H. anion exchange no. 2 100 mM NaCl eluate.

FIG. 7. Toyopearl 650 M (phenyl) HIC chromatograph. Buffer A: 6 M urea, 25 mM Tris-HCl, pH 7.5, 1 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>; Buffer B: 6 M urea, 25 mM Tris-HCl, pH 7.5; Gradient: Step, 100% B for the flow-through and 0% B for strip; and Flow rate: 10 ml/min.

FIG. 8. SDS-PAGE analysis troponin lot after hydrophobic interaction chromatography in 16 $\times$  tris-glycine gel, under nonreducing conditions. A-H refer to lanes in the SDS-PAGE gel. A. Sulfitolyzed troponin Lot 3L4 standard; B. AEX step no. 2, troponin eluate pool; C. HIC load (w/1M ammonium sulfate); D. HIC flowthrough (troponin product); E. HIC low salt eluate (column strip); F. lot 3L5 sulfitolyzed troponin product.

FIG. 9. Quantitation of rTnI on Zorbax C3.

FIG. 10. Troponin I LysC mapping.

FIG. 11. 3D S-PAGE analysis of sulfitolyzed troponin reduction with dithiothreitol for 45 mins. at ambient temperature. One ng/ml TnI in 6 M urea, 25 mM tris, 0.15 M NaCl pH 7.5, run on 16 $\times$  tris-glycine gel. 1. 10 $\times$  Mark 12 MW Stds; 2. 9 $\times$  sulfitolyzed TnI; 3. 0.55 mM DTT; 4. 0.10 mM DTT; 5. 0.2 mM DTT; 6. 0.3 mM DTT; 7. 0.5 mM DTT; 8. 1.0 mM DTT.

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TITLE: PURIFICATION OF HUMAN TROPONIN I

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# ABSTRACT OF THE INVENTION

FIG. 1A and 1B. A. Proposed reaction for oxidative sulfitolysis. B. Cleanup of disulfide bond by sodium sulfite to form the Ssulfide derivative.

FIG. 2. Preparation and washing of TnI-containing inclusion bodies.

FIG. 3. Summary of rTroponin-I preparation.

FIG. 4. Q-Sepharose FF chromatography of Troponin I. Buffer A: 6 M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6 M urea, 25 mM Tris-HCl, pH 7.5, 20 mM NaCl; Gradient: Step, 100% B for the flowthrough and 0% B for the strip; and Flow rate: 10 ml/min.

FIG. 5. 30 ml Q-sepharose FF ion exchange chromatography. Buffer A: 6M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6M urea, 25 mM Tris-HCl, pH 7.5, 2M NaCl; Gradient: Step, 40 B for elution and 50 B for strip; and Flow rate: 20 ml/min.  
 FIG. 6. SDS-PAGE analysis troponin I after anion exchange steps no. 1 and no. 2 in 16 tris-glycine gel, under nonreducing conditions. A-H refer to lanes in the SDS-PAGE gel. A. Sulfite-lyzed troponin I; B. solubilized inclusion bodies; C. sulfite-lyzed inclusion bodies (AEX No. 1 load); D. anion exchange no. 1 flowthrough; E. anion exchange no. 1 salt eluate; F. anion exchange no. 2 load; G. anion exchange no. 2 flowthrough; and, H. anion exchange no. 2 100 mM NaCl eluate.

FIG. 7. Toyopearl 650M (phenyl) HIC chromatography. Buffer A: 6M urea, 25 mM Tris-HCl, pH 7.5, 1M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>; Buffer B: 6M urea, 25 mM Tris-HCl, pH 7.5; Gradient: Step, 100 B for the flowthrough and 0 B for strip; and Flow rate: 10 ml/min.

FIG. 8. SDS-PAGE analysis troponin I after hydrophobic interaction chromatography in 16 tris-glycine gel, under nonreducing conditions. A-F refers to lanes in the SDS-PAGE gel. A. Sulfite-lyzed troponin I; B. AEX step no. 2, troponin eluate pool; C. HIC load (w/1M ammonium sulfate); D. HIC flowthrough (troponin product); E. HIC low salt eluate (column strip); F. lot SDS sulfite-lyzed troponin product.

FIG. 9. Quantitation of rTnI on Zorbax C3.

FIG. 10. Troponin I LysC mapping.

FIG. 11. SDS-PAGE analysis of sulfite-lyzed troponin reduction with dithiothreitol for 45 mins. at ambient temperature. One mg/ml TnI in 6M urea, 25 mM Tris, 0.1M NaCl pH 7.5, run on 16 tris-glycine gel. 1.10, Mark 12 MW Stds; 2.9, sulfite-lyzed TnI; 3. 1.0 mM DTT; 4. 0.5 mM DTT; 5. 0.1 mM DTT; 6. 0.3 mM DTT; 7. 0.5 mM DTT; 8. 1.0 mM DTT.

AB The invention is directed to methods for purifying **Troponin I**, particularly recombinant Troponin I produced in a bacterial expression system. Recombinant Troponin I can be advantageously purified after reversibly protecting the free sulfhydryl groups, e.g., by forming sulfates. In a specific example, Troponin I reacted with sodium tetrafluoroborate yielded sulfite-lyzed Troponin I, which was purified by chromatography on an anion exchanger, followed by hydrophobic interaction chromatography. Facile deprotection of the sulfhydryl groups yields a highly purified product ready for refolding.

CLMN 20-11 Figures).

FIGS. 1A and 1B. A. Proposed reaction for oxidative sulfiteolysis. B. Cleavage of disulfide bond by sodium sulfite to form the Ssulfo derivative.

FIG. 2. Preparation and washing of TnI-containing inclusion bodies.

FIG. 3. Summary of rTroponin-I preparation.

FIG. 4. Q-sepharose FF chromatography of **Troponin I**.

Buffer A: 6M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6M urea, 25 mM Tris-HCl, pH 7.5, 2M NaCl; Gradient: Step, 40 B for elution and 50 B for strip; and Flow rate: 150 ml/min.

FIG. 5. 300 ml Q-sepharose FF chromatography. Buffer A: 6M urea, 25 mM Tris-HCl, pH 7.5, 100 mM; Buffer B: 6M urea, 25 mM Tris-HCl, pH 7.5, 2M NaCl; Gradient: Step, 40 B for elution and 50 B for strip; and Flow rate: 20 ml/min.

FIG. 6. SDS-PAGE analysis troponin I after anion exchange steps no. 1 and no. 2 in 16 tris-glycine gel, under nonreducing conditions. A-H refer to lanes in the SDS-PAGE gel. A. Sulfite-lyzed troponin I; B. solubilized inclusion bodies; C. sulfite-lyzed inclusion bodies (AEX No. 1 load); D. anion exchange no. 1 flowthrough; E. anion exchange no. 1 salt eluate; F. anion exchange no. 2 load; G. anion exchange no. 2 flowthrough; and, H. anion exchange no. 2 100 mM NaCl eluate.

FIG. 7. Toyopearl 650M (phenyl) HIC chromatography. Buffer A: 6M urea, 25 mM Tris-HCl, pH 7.5, 1M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>; Buffer B: 6M urea, 25 mM Tris-HCl, pH 7.5; Gradient: Step, 100 B for the flowthrough and 0 B for strip; and Flow rate: 10 ml/min.

FIG. 8. SDS-PAGE analysis troponin I after hydrophobic interaction chromatography in 16 tris-glycine gel, under nonreducing conditions. A-F refers to lanes in the SDS-PAGE gel. A. Sulfite-lyzed troponin I; B. AEX step no. 2, troponin eluate pool; C. HIC load (w/1M ammonium sulfate); D. HIC flowthrough (troponin product); E. HIC low salt

eluate (column strip); F. lot 3L5 sulfiteylzed troponin product.

FIG. 9. Quantitation of rTnI on Zorbax 63.

FIG. 10. **Troponin I** lysC mapping.

FIG. 11. SDS-PAGE analysis of sulfiteylzed troponin reduction with dithiothreitol for 45 mins. at ambient temperature. One mg/ml TnI in 6M urea, 25 mM tris, 0.15M NaCl pH 7.5, run on 16% tris-glycine gel. 1.16%, Mark 12 MW Stds; 2.9., sulfiteylzed TnI; 3. 0.05 mM DTT; 4. 0.10 mM DTT; 5. 0.2 mM DTT; 6. 0.3 mM DTT; 7. 0.5 mM DTT; 8. 1.0 mM DTT.